### AIMS AND SCOPE

Although total synthesis reached extraordinary levels of sophistication in the last century, the development of practical and efficient synthetic methodologies is still in its infancy. Achieving chemical reactions that are highly selective, economical, safe, resource- and energy-efficient, and environmentally benign is a primary challenge to chemistry in this century. Realizing this goal will demand the highest level of scientific creativity, insight and understanding in a combined effort by academic, government and industrial chemists and engineers.

Advanced Synthesis & Catalysis promotes that process by publishing high-impact research results reporting the development and application of efficient synthetic methodologies and strategies for organic targets that range from pharmaceuticals to organic materials. Homogeneous catalysis, biocatalysis, organocatalysis and heterogeneous catalysis directed towards organic synthesis are playing an ever increasing role in achieving synthetic efficiency. Asymmetric catalysis remains a topic of central importance. In addition, Advanced Synthesis & Catalysis includes other areas that are making a contribution to green synthesis, such as synthesis design, reaction techniques, flow chemistry and continuous processing, multiphase catalysis, green solvents, catalyst immobilization and recycling, separation science and process development.

Practical processes involve development of effective integrated strategies, from an elegant synthetic route based on mechanistic and structural insights at the molecular level through to process optimization at larger scales. These endeavors often entail a multidisciplinary approach that spans the broad fields chemistry, biology, and engineering and involve contributions from academic, government, and industrial laboratories.

The unique focus of *Advanced Synthesis & Catalysis* has rapidly made it a leading organic chemistry and catalysis journal. The goal of *Advanced Synthesis & Catalysis* is to help inspire a new era of chemical science, based on the efforts of synthetic chemists and on interdisciplinary collaboration, so that chemistry will make an even greater contribution to the quality of life than it does now.



succeeding Journal für praktische Chemie (founded in 1828)

ASC
2-Year Impact Factor 2008
5.619
A Record High
for Organic Chemistry

2009, 351, 17, Pages 2733-3012

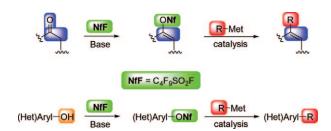
Issue 16/2009 was published online on November 5, 2009

### REVIEWS

Nine Times Fluoride can be Good for your Syntheses. Not just Cheaper: Nonafluorobutanesulfonates as Intermediates for Transition Metal-Catalyzed Reactions

Adv. Synth. Catal. 2009, 351, 2747-2763

Jens Högermeier, Hans-Ulrich Reissig\*



\*\*WILEY InterScience\*

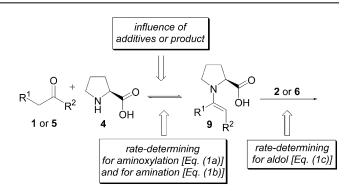
2747

### **COMMUNICATIONS**

### 2765 A Coherent Mechanistic Rationale for Additive Effects and Autoinductive Behaviour in Proline-Mediated Reactions

Adv. Synth. Catal. 2009, 351, 2765-2769

Natalia Zotova, Antonio Moran, Alan Armstrong,\* Donna G. Blackmond\*



# 2770 Gold- and Silver-Catalyzed Intramolecular Hydroamination of Terminal Alkynes: Water-Triggered Chemo- and Regioselective Synthesis of Fused Tricyclic Xanthines

Adv. Synth. Catal. 2009, 351, 2770-2778

Deju Ye, Xu Zhang, Yu Zhou, Dengyou Zhang, Lei Zhang, Hengshuai Wang, Hualiang Jiang, Hong Liu\*

## **2779** DiPAMPs Big Brother "*i*-Pr-SMS-Phos" Exhibits Exceptional Features Enhancing Rhodium(I)-Catalyzed Hydrogenation of Olefins

Adv. Synth. Catal. 2009, 351, 2779-2786

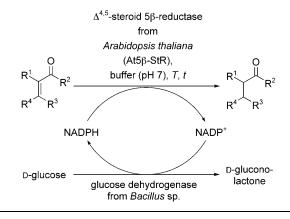
Michel Stephan,\* Damjan Šterk, Barbara Mohar

*i*-Pr-SMS-Phos ligand displayed a boosted catalyst activity and enantioselectivity in the rhodium(I)-catalyzed hydrogenation of a wide-range of dehydro- $\alpha$ -amido acids, itaconates, acrylates, enamides, enol acetates and  $\alpha$ , $\alpha$ -diarylethylenes.

### 2787 Recombinant Δ<sup>4,5</sup>-Steroid 5β-Reductases as Biocatalysts for the Reduction of Activated C=C-Double Bonds in Monocyclic and Acyclic Molecules

Adv. Synth. Catal. 2009, 351, 2787-2790

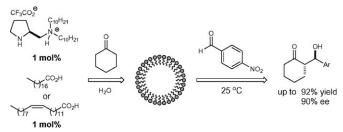
Edyta Burda, Marina Kraußer, Gabriele Fischer, Werner Hummel, Frieder Müller-Uri, Wolfgang Kreis,\* Harald Gröger\*



## 2791 Effect of Long Chain Fatty Acids on Organocatalytic Aqueous Direct Aldol Reactions

Adv. Synth. Catal. 2009, 351, 2791-2796

Nobuyuki Mase,\* Naoyasu Noshiro, Asuka Mokuya, Kunihiko Takabe



Organocatalytic Asymmetric Synthesis of Protected  $\alpha,\beta$ -Diamino Acids

Adv. Synth. Catal. 2009, 351, 2797-2800

Zugui Shi, Peiyuan Yu, Pei Juan Chua, Guofu Zhong\*

R = aromatic, heteroaromatic, aliphatic up to 99% yield, >99% ee, 97:3 dr

Highly Efficient Ion-Tagged Catalyst for the Enantioselective Michael Addition of Aldehydes to Nitroalkenes

Adv. Synth. Catal. 2009, 351, 2801-2806

Marco Lombardo,\* Michel Chiarucci, Arianna Quintavalla, Claudio Trombini

Ph  
Ph  
Ph  

$$R^2$$
 $R^3$ 
 $NO_2$ 
 $R^3$ 
 $NO_2$ 
 $R^3$ 
 $NO_2$ 
 $R^3$ 
 $NO_2$ 
 $R^3$ 
 $NO_2$ 
 $R^3$ 
 $NO_2$ 
 $R^3$ 
 $NO_2$ 

Y up to >99%, syn ee up to >99.5%, syn:anti up to 98:2

Development of a New Spiro-BOX Ligand and Its Application in Highly Enantioselective Palladium-Catalyzed Cyclization of 2-Iodoanilines with Allenes

Adv. Synth. Catal. 2009, 351, 2807-2810

☐ Wei Shu, Qiong Yu, Shengming Ma\*

Highly Enantioselective Michael Addition of  $\alpha$ -Substituted Cyano Ketones to  $\beta$ , $\gamma$ -Unsaturated  $\alpha$ -Keto Esters using Bifunctional Thiourea-Tertiary Amine Catalysts: An Easy Access to Chiral Dihydropyrans

Adv. Synth. Catal. 2009, 351, 2811-2816

Sheng-Li Zhao, Chang-Wu Zheng, Hai-Feng Wang, Gang Zhao\*

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2807

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2817 Enantioselective Electrophilic Amination of  $\alpha$ Cyanothioacetates with Azodicarboxylates Catalyzed by an Axially Chiral Guanidine Base

Adv. Synth. Catal. 2009, 351, 2817-2821

- Masahiro Terada,\* Daisuke Tsushima, Megumi Nakano
- Boc NC SEt

  NH2

  Boc NC SEt

  R (R)-Cat. (0.1 1 mol%) (R)-Cat. (0.1 1 mol%)Boc NH R

  Boc NH R

  up to 99% ee
- **2822** Enantioselective Synthesis of Dihydrocoumarins *via* N-Heterocyclic Carbene-Catalyzed Cycloaddition of Ketenes and *o*-Quinone Methides

Adv. Synth. Catal. 2009, 351, 2822-2826

Hui Lv, Lin You, Song Ye\*

- Ar C=O Ph ... N Ph Ph OTBS (10 mol%)

  DME, MeOH (0.6 equiv.)

  Cis:trans up to 9:1 ee up to 99% (cis)
- **2827** Palladium-Catalyzed Decarboxylative *sp-sp*<sup>2</sup> Cross-Coupling Reactions of Aryl and Vinyl Halides and Triflates with α,β-Ynoic Acids using Silver Oxide

Adv. Synth. Catal. 2009, 351, 2827-2832

☐ Hyunseok Kim, Phil Ho Lee\*

X = Br, I, OTf

FG = Me, n-Bu, MeO, CHO, Ac, EtO<sub>2</sub>C, CN, Ph; FG-C<sub>6</sub>H<sub>4</sub> = 2-pyridyl, 2-thienyl R = n-Bu, Ph, 3-Me-C<sub>6</sub>H<sub>4</sub>, 3-MeO-C<sub>6</sub>H<sub>4</sub>, 4-MeO-C<sub>6</sub>H<sub>4</sub>

**2833** Facile Synthesis of Polycyclic Fluorene Derivatives *via* a Palladium-Catalyzed Coupling, Propargyl-Allenyl Isomerization and Schmittel Cyclization Sequence

Adv. Synth. Catal. 2009, 351, 2833-2838

Ruwei Shen, Linzhu Chen, Xian Huang\*

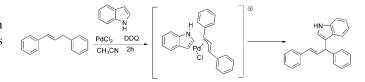
- $\mathsf{EWG} \underbrace{\begin{array}{c} \mathsf{EWG} \\ \mathsf{Pd}(\mathsf{Ph}_{\mathsf{P}}\mathsf{P}_{\mathsf{2}}\mathsf{Cl}_{\mathsf{2}} \\ \mathsf{Cul} \\ \mathsf{Toluene}, \mathsf{Et}_{\mathsf{3}}\mathsf{N} \\ \mathsf{R}^{\mathsf{3}} \end{array}}_{\mathsf{R}^{\mathsf{3}}} \underbrace{\begin{array}{c} \mathsf{R}^{\mathsf{3}} \\ \mathsf{R}^{\mathsf{3}}$
- **2839** Intramolecular Photochemical Cross-Coupling Reactions of 3-Acyl-2-haloindoles and 2-Chloropyrrole-3-carbaldehydes with Substituted Benzenes

Adv. Synth. Catal. 2009, 351, 2839-2844

- Shen-Ci Lu, Xi-Xia Zhang, Zong-Jun Shi, Yu-Wei Ren, Bing Li, Wei Zhang\*
- $R^1$   $R^2$   $R^2$
- **2845** Efficient Palladium-Catalyzed Oxidative Indolation of Allylic Compounds with DDQ *via sp*<sup>3</sup> C–H Bond Activation and Carbon-Carbon Bond Formation Under Mild Conditions

Adv. Synth. Catal. 2009, 351, 2845-2849

☐ Hanjie Mo, Weiliang Bao\*



2855

2887

2897

Iridium-Catalyzed Selective Synthesis of 4-Substituted Benzofurans and Indoles via Directed Cyclodehydration

Adv. Synth. Catal. 2009, 351, 2850-2854

Kyoji Tsuchikama, Yu-ki Hashimoto, Kohei Endo, Takanori Shibata\*

### **FULL PAPERS**

Gold Catalysis: Anellated Heterocycles and Dependency of the Reaction Pathway on the Tether Length

Adv. Synth. Catal. 2009, 351, 2855-2875

🖳 A. Stephen K. Hashmi,\* Sreekumar Pankajakshan, Matthias Rudolph, Elisabeth Enns, Thomas Bander, Frank Rominger, Wolfgang Frey

Reusable Gold(I) Catalysts with Unique Regioselectivity for Intermolecular Hydroamination of Alkynes

Adv. Synth. Catal. 2009, 351, 2876-2886

Antonio Leyva, Avelino Corma\*

Semiconductor-Gold Nanocomposite Catalysts for the Efficient Three-Component Coupling of Aldehyde, Amine and Alkyne in Water

Adv. Synth. Catal. 2009, 351, 2887-2896 Leng Leng Chng, Jun Yang, Yifeng Wei, Jackie Y. Ying\*

Axially Chiral Phosphine-Oxazoline Ligands in Silver(I)-Catalyzed Asymmetric Mannich Reaction of Aldimines with Trimethylsiloxyfuran

Adv. Synth. Catal. 2009, 351, 2897-2902

☐ Hong-Ping Deng, Yin Wei, Min Shi\*

 $R^1$  = styryl;  $R^2$  = phenyl or solvent-free r.t. or 80 °C, 24 h regioselectivity up to 90% PbS-Au  $R^1$ -CHO +  $R^2R^3NH$  + H<sub>2</sub>O, 100 °C

12 - 13 h up to 95% yield

CH2Cl2, CH3CN

R3NH2

Synthesis of Selectively Mono-N-Arylated Aliphatic Diamines via Iridium-Catalyzed Amine Alkylation

Adv. Synth. Catal. 2009, 351, 2903-2911

Benoît Blank, Stefan Michlik, Rhett Kempe\*

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2912 Expanded Heterogeneous Suzuki–Miyaura Coupling Reactions of Aryl and Heteroaryl Chlorides under Mild Conditions

Adv. Synth. Catal. 2009, 351, 2912-2920

- Dong-Hwan Lee, Minkee Choi, Byung-Woo Yu, Ryong Ryoo,\* Abu Taher, Shahin Hossain, Myung-Jong Jin\*
- Ar-CI  $+ (HO)_{2}B-Ar'$   $+ (HO)_{2}B-Ar'$  + (H
- 2921 Novel Enantioselective Sequentially Rhodium(I)/BINAP-Catalyzed Cycloisomerization–Hydrogenation–Isomerization– Acetalization (CIHIA)

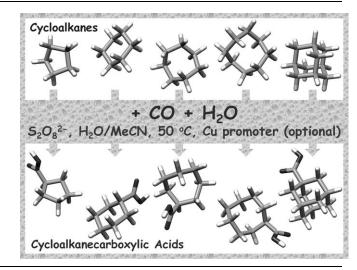
Adv. Synth. Catal. 2009, 351, 2921-2935

- Nadine Körber, Frank Rominger, Thomas J. J. Müller\*
- One-Pot

  | R | [RhCl(cod)]<sub>2</sub>, (R)-BINAP | H | R | AgBF<sub>4</sub> | DCE, r.t. |
  | OH | then: H<sub>2</sub> (5 bar) | H | (R,R,R)-4 (34 70%; 12 examples) | ee > 99%)
- 2936 Metal-Free and Copper-Promoted Single-Pot Hydrocarboxylation of Cycloalkanes to Carboxylic Acids in Aqueous Medium

Adv. Synth. Catal. 2009, 351, 2936-2948

Marina V. Kirillova, Alexander M. Kirillov, Armando J. L. Pombeiro\*



**2949** Copper-Catalyzed *N*-Alkylation of Sulfonamides with Benzylic Alcohols: Catalysis and Mechanistic Studies

Adv. Synth. Catal. 2009, 351, 2949-2958

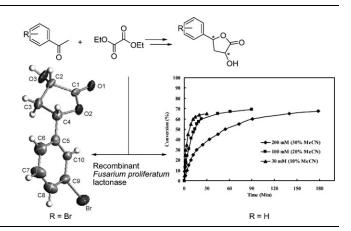
- Xinjiang Cui, Feng Shi,\* Man Kin Tse, Dirk Gördes, Kerstin Thurow, Matthias Beller, Youquan Deng\*
- O, O NH<sub>2</sub> + OH 1 mol% [Cu] 150 °C, 12 h 40-mL tube

29 samples up to 99% isolated yield

**2959** Facile Synthesis of Enantiopure 4-Substituted 2-Hydroxy-4-butyrolactones using a Robust *Fusarium* Lactonase

Adv. Synth. Catal. 2009, 351, 2959-2966

Bing Chen, Hai-Feng Yin, Zhen-Sheng Wang, Jian-He Xu,\* Li-Qiang Fan, Jian Zhao



Activated α,β-Unsaturated Aldehydes as Substrate of Dihydroxyacetone Phosphate (DHAP)-Dependent Aldolases in the Context of a Multienzyme System

Adv. Synth. Catal. 2009, 351, 2967-2975

Israel Sánchez-Moreno, Laura Iturrate, Elisa G. Doyagüez, Juan Antonio Martínez, Alfonso Fernández-Mayoralas, Eduardo García-Junceda\*

Asymmetric Reduction of Activated Alkenes by Pentaerythritol Tetranitrate Reductase: Specificity and Control of Stereochemical Outcome by Reaction Optimisation

Adv. Synth. Catal. 2009, 351, 2976-2990

Anna Fryszkowska, Helen Toogood, Michiyo Sakuma, John M. Gardiner, Gill M. Stephens,\* Nigel S. Scrutton\*

The Addition Reaction of Samarium Enolates and 2-Haloenolates Derived from Esters, and Amides to Imines. Totally Stereoselective Synthesis of Enantiopure 3,4-Diamino Esters or Amides

Adv. Synth. Catal. 2009, 351, 2991-3000

☐ José M. Concellón,\* Humberto Rodríguez-Solla, Carmen Simal, Vicente del Amo, Santiago García-Granda, M. Rosario Díaz

N-Heterocyclic Carbene-Mediated Enantioselective Addition of Phenols to Unsymmetrical Alkylarylketenes

Adv. Synth. Catal. 2009, 351, 3001-3009

Carmen Concellón, Nicolas Duguet, Andrew D. Smith\*

Supporting information on the WWW (see article for access details)

\*Author to whom correspondence should be addressed.

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